

17. (Twice Amended) A method for producing an antimycobacterial compound of the

formula:

wherein R₁ is H; and

wherein R2 is phenyl, substituted phenyls, napthyls and substituted napthyls or

wherein R_1R_2 = optionally substituted carbocyclic groups;

which comprises:

refluxing

CONHNH₂

(1)

with absolute ethanol to produce a solution;

adding a carbonyl compound comprising the formula of:

 R_3COR_4 (2)

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wherein R₃ = H or CH₃; and

Soft By wherein $R_4 = C_1$ to C_{14} alkyl, C_2 to C_{10} substituted alkyl, C_2 to C_{10} alkenyl, C_2 to C_9 substituted alkenyl, C_3 to C_7 cycloalkyl, C_3 to C_7 substituted cycloalkyl, phonyl, substituted phonyl, C_7 to C_{16} phonylalkyl, C_7 to C_{16} substituted phonylalkyl, behazyl, substituted benzyl, naphthyl, substituted naphthyl, heterocycle, substituted heterocycle, hald hydroxy, amino, or carboxy; or

wherein $R_3 R_4 = C_4$ to C_8 cycloalkyl or C_4 to C_{10} substituted cycloalkyl;

to the solution to produce a reaction mixture, the reaction mixture having a mole ratio of carbonyl compound to compound (1) of 1.67 to 1.00;

distilling the reaction mixture;

precipitating a solid from the reaction mixture:

filtering the solid; and

drying the solid to obtain I.

Please add the following claims:

24. (New) The method of claim 17 wherein R₂ of compound I is phenyl substituted with 1 to 3 substituents selected from the group consisting of a halogen, a hydroxyl, a methoxy, a benzyloxy, a phenoxy, a trifluoromethyl, an isopropyl, and a thiomethyl group.

25. (New) The method of claim 24 wherein R_2 of compound I = 4-iso- $C_3H_7C_6H_4$, 2,5-di(Cl)C₆H₃, 2,3,5-tri(F)C₆H₂, 2-F-4-CF₃C₆H₃, 3,4,5-tri(F)C₆H₂, 2-Cl-6-CH₃O-iso-C₉H₄N, 2-F-3-Cl-6-CF₃C₆H₂, 2,4-di(CF₃)C₆H₃, 2,6-di(F)-3-Cl-C₆H₂, 2-F-3-Cl-5-CF₃-C₆H₂, 2-F-5-Br-C₆H₃, 2-CH₃S-C₆H₄, 2-O-C₇H₇C₆H₄, 3-O-C₇H₇C₆H₄, 4-O-C₇H₇C₆H₄, 2,4,5-tri(F)C₆H₂, 2-F-5-I-C₆H₃, 2,3,4-tri(OH)C₆H₂, 4-C₆H₄-CH=NNHCO-4-C₅H₄N, 4-C₆H₄-O-CH₂CH₂CH₂CH₃, 4-C₆H₄NO₂, 2-C₆H₄OH, 4-OH-3-OCH₃C₆H₃, 4-C₆H₄OCH₃, 3-C₆H₄OCH₃, 4-C₆H₄F, 3,5-di(CH₃)-4-O-C₇H₇, 2-C₆H₄OH, 4-OH-3-OCH₃C₆H₃, 4-C₆H₄OCH₃, 4-C₆H₄OCH₃, 4-C₆H₄OCH₃, 4-C₆H₄F, 3,5-di(CH₃)-4-O-C₇H₇, 2-C₆H₄OCH₃, 4-C₆H₄OCH₃, 4-C₆H₄

 $F-4-OCH_3C_6H_3,\ 2-ClC_6H_4,\ 4-BrC_6H_4,\ 3-C_6H_4NO_2,\ 4-C_6H_4O(CH_2)_5CH_3,\ 2-Cl-5-NO_2C_6H_3,\ 4-Cl-3-NO_2C_6H_3,\ 2-C_6H_4NO_2,\ 2-6-di(Cl)C_6H_3,\ 2,3-di(Cl)C_6H_3,\ 3,4-di(F)C_6H_3,\ 2,6-di(F)C_6H_3,\ 3,4-di(Cl)C_6H_3,\ 0-C_6H_4Cl.$

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26. (New) The method of claim 17 wherein R_2 of compound I =

or

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27. (New) The method of claim 17 wherein R_1R_2 of compound I is

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